Superconductivity induced by phosphorus doping and its coexistence with ferromagnetism in EuFe$_2$(As$_{0.7}$P$_{0.3}$)$_2$

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We have studied EuFe$_2$(As$_{0.7}$P$_{0.3}$)$_2$ by the measurements of x-ray diffraction, electrical resistivity, thermopower, magnetic susceptibility, magnetoresistance and specific heat. Partial substitution of As with P results in the shrinkage of lattice, which generates chemical pressure to the system. It is found that EuFe$_2$(As$_{0.7}$P$_{0.3}$)$_2$ undergoes a superconducting transition at 26 K, followed by ferromagnetic ordering of Eu^{2+} moments at 20 K. This finding is the first observation of superconductivity stabilized by internal chemical pressure, and supplies a rare example showing coexistence of superconductivity and ferromagnetism in the ferro-arsenide family.

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Recently, high-temperature superconductivity has been discovered in a family of materials containing FeAs layers.[1, 2, 3, 4] The superconductivity is induced by doping charge carriers into a parent compound, characterized by an antiferromagnetic (AFM) spin-density-wave (SDW) transition associated with the FeAs layers.[5, 6] Electron doping has been realized by the partial substitutions of F-for-O,[1] vacancy-for-O,[3] Th-for-Ln,[4] Co/Ni-for-Fe,[7, 8, 9, 10]. Examples of hole doping include the partial substitutions of Sr-for-Ln[11] and K-for-Ba/Sr/Eu[12, 13, 14]. All the chemical doping suppresses the long-range SDW order, eventually resulting in the emergence of superconductivity. Up to now, no superconductivity has been reported through doping at the As site. Apart from carrier doping, application of hydrostatic pressure is also capable of stabilizing superconductivity.[15, 16, 17]

Among the parent compounds in ferro-arsenide family, EuFe$_2$As$_2$ exhibits peculiar behavior because the moments of Eu^{2+} ions order antiferromagnetically at relatively high temperature of 20 K.[18, 19, 20] Magnetoresistance measurements on EuFe$_2$As$_2$ crystals[21] suggest a strong coupling between the magnetism of Eu^{2+} ions and conduction electrons, which may affect or even destroy superconductivity. For example, though Ni doping in BaFe$_2$As$_2$ leads to superconductivity up to 21 K,[10] ferromagnetism rather than superconductivity was found in EuFe$_2$As$_2$ by a systematic Ni doping[22]. Another relevant example is that the superconducting transition temperature of (Eu,K)Fe$_2$As$_2$[14] is 32 K, substantially lower than those of (Ba,K)Fe$_2$As$_2$ (T_c=38 K)[12] and (Sr,K)Fe$_2$As$_2$ (T_c=37 K)[13]. Resistivity measurement under hydrostatic pressures[17] on EuFe$_2$As$_2$ crystals showed a resistivity drop at 29.5 K. However, no zero

resistivity could be achieved, which was ascribed to the AFM ordering of the Eu^{2+} moments[17].

While hetero-valent substitution generally induces charge carriers, iso-valent substitution may supply chemical pressure. The latter substitution is of particular interest in EuFe$_2$As$_2$ when As is partially replaced by P, as suggested theoretically in order to search for the magnetic quantum criticality without changing the number of Fe 3d-electrons[23]. In this Letter, we demonstrate bulk superconductivity at 26 K in EuFe$_2$(As$_{0.7}$P$_{0.3}$)$_2$. For the first time, superconductivity has been realized through the doping at the As site in the iron arsenide system. Strikingly, we observe coexistence of ferromagnetic ordering of Eu^{2+} moments with superconductivity below 20 K.

Polycrystalline samples of EuFe$_2$(As$_{0.7}$P$_{0.3}$)$_2$ were synthesized by solid state reaction with EuAs, Fe$_2$As and Fe$_2$P. EuAs was presynthesized by reacting Eu grains and As powders at 873 K for 10 h, then 1073 K for 10 h and finally 1223 K for another 10 h. Fe$_2$As was prepared by reacting Fe powers and As powders at 873 K for 10 h then 1173 K for 0.5 h. Fe$_2$P was presynthesized by heating Fe powders and P powders very slowly to 873 K and holding for 10 h. Powders of EuAs, Fe$_2$As and Fe$_2$P were weighed according to the stoichiometric ratio, ground and pressed into pellets in an argon-filled glove-box. The pellets were sealed in evacuated quartz tubes and annealed at 1273 K for 20 h then cooled slowly to room temperature. The phase purity of the samples was investigated by powder X-ray diffraction, using a D/Max-rA diffractometer with Cu-K$_\alpha$ radiation and a graphite monochromator. The XRD data were collected with a step-scan mode in the 2θ range from 10° to 120°. The structural refinement was performed using the programme RIETAN 2000.[24]

The electrical resistivity was measured on bar-shaped samples using a standard four-probe method. The applied current density was $\sim$ 0.5 A/cm^2. The measurements of magnetoresistance, specific heat, ac magnetic susceptibility and thermopower were performed on a
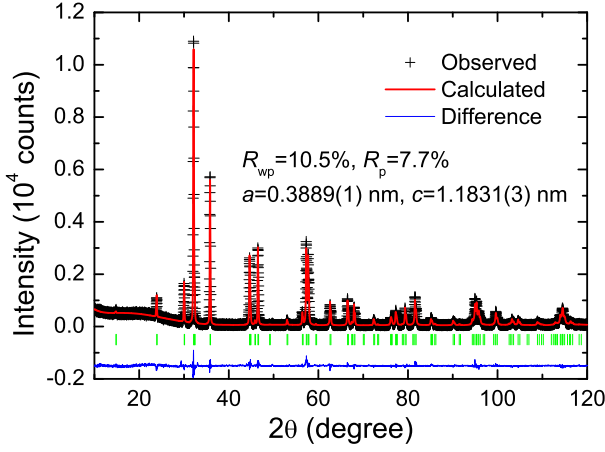
FIG. 1: (color online). X-ray powder diffraction pattern at room temperature and the Rietveld refinement profile for $EuFe_2(As_{0.7}P_{0.3})_2$.

Quantum Design Physical Property Measurement System (PPMS-9). DC magnetic properties were measured on a Quantum Design Magnetic Property Measurement System (MPMS-5).

Figure 1 show the XRD pattern of $EuFe_2(As_{0.7}P_{0.3})_2$ at room temperature, together with the profile of the Rietveld refinement using the space group of $I4/mmm$. No additional diffraction peak is observed. The refined lattice parameters are a=3.889(1)Å and c=11.831(3)Å. Compared with those of the undoped $EuFe_2As_2$[19], the a-axis is decreased by 0.35%, the c-axis is shortened by 1.8% and the cell volume shrinks by 3.2% for $EuFe_2(As_{0.7}P_{0.3})_2$. These results suggest that the isovalent substitution of As with P indeed generates chemical pressure to the system. In addition, the As(P) position z decreases, indicating that the As(P) atoms move toward the Fe planes. As a consequence, the Fe-As(P)-Fe angle increases from 110.15° to 111.48°.

Figure 2(a) shows the temperature dependence of resistivity (ρ) for $EuFe_2(As_{0.7}P_{0.3})_2$ under zero field. The anomaly associated with the SDW transition in undoped $EuFe_2As_2$[19] is completely suppressed. The resistivity is linear with temperature down to $\sim$ 90 K and shows upward deviation from the linearity at lower temperatures. The resistivity ratio of $R(300K)/R(30K)$ is 5.2, indicating high quality of the present sample. Below 29 K, the resistivity drops steeply, suggesting a superconducting transition. The midpoint of the transition is 26 K. On closer examination shown in the inset of Fig. 1(a), however, a small resistivity peak is observed around 16 K, which coincides with the ferromagnetic ordering of Eu^{2+} moments (to be shown below). This observation is reminiscent of the reentrant superconducting behavior as observed, for example, in $RNi_2B_2C_2$ (R=Tm, Er, Ho)[25].

Figure 2(b) shows the temperature dependence of ther-

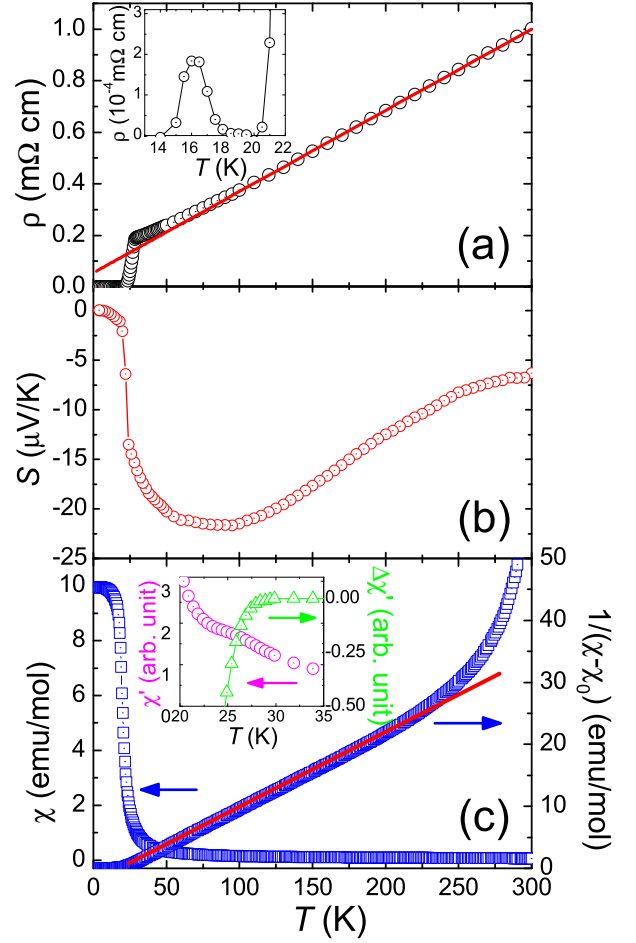
FIG. 2: (color online). Temperature dependence of resistivity (a), thermopower (b), and magnetic susceptibility (c) for the $EuFe_2(As_{0.7}P_{0.3})_2$ sample. The inset of (a) shows an expanded plot. The inset of (c) shows the real part of ac susceptibility χ' (left axis) for the same sample. A diamagnetic signal (right axis) is obtained after subtraction of paramagnetic contribution of Eu^{2+} moments.

mopower (S). The thermopower in the whole temperature range is negative, indicating that the dominant charge carriers are electron-like. $|S|$ decreases sharply below 29 K, corresponding to the superconducting transition. However, the $|S|$ value does not drop to zero until $\sim$ 13 K, in relation with the ferromagnetic ordering of the Eu^{2+} moments.

In figure 2(c), we show the temperature dependence of field-cooling dc magnetic susceptibility for $EuFe_2(As_{0.7}P_{0.3})_2$ under μ_0H_{dc}=0.1 T. The obvious deviation of linearity in χ^{-1} above 230 K is probably due to the presence of trace amount of ferromagnetic Fe_2P impurity[26]. The χ data between 50 K and 200 K can be well described by the modified Curie-Weiss law,

$$\chi = \chi_0 + \frac{C}{T - \theta}, \qquad (1)$$

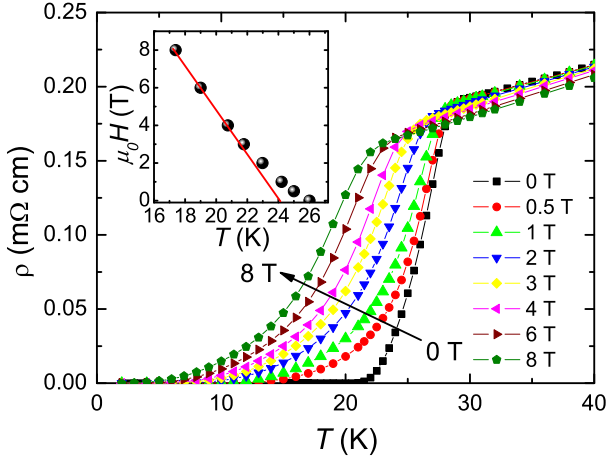where χ_0 denotes the temperature-independent term, C

FIG. 3: (color online). Field dependence of resistive transition for $EuFe_2(As_{0.7}P_{0.3})_2$ sample. The inset shows the upper critical fields as a function of temperature.
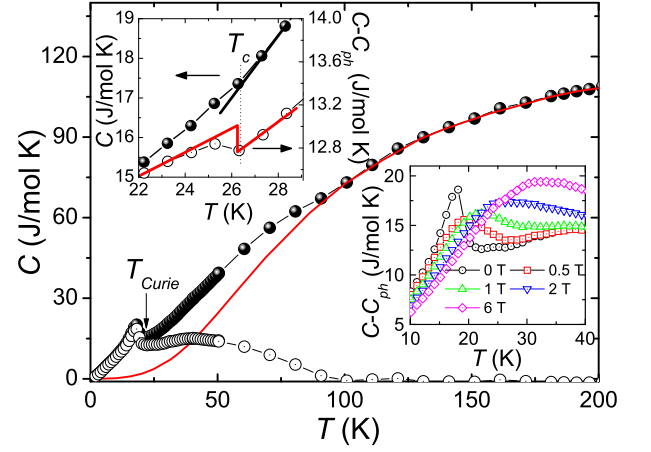

FIG. 4: (color online). Temperature dependence of specific heat before (solid symbol) and after (open symbol) deduction of lattice contribution for the $EuFe_2(As_{0.7}P_{0.3})_2$ sample. The solid curve represents the lattice contribution fitted by the Debye model. See text for details. The upper left panel shows the specific heat anomaly around 26 K, which is ascribed to the superconducting transition. The lower right panel shows the magnetic specific heat anomaly around 20 K under magnetic fields.

the Curie-Weiss constant and θ the Weiss temperature. The fitting yields C= 8.1(1) emu·K/mol and θ=22(1) K. The corresponding effective moment P_{eff}=8.0(1) μ_B per formula unit, close to the theoretical value of 7.94 μ_B for a free Eu^{2+} ion. χ increases steeply with decreasing temperature below 20 K and becomes gradually saturated. Field dependence of magnetization gives a saturated magnetic moment of $6.9(1)\mu_B$/f.u., as expected for fully paralleled Eu^{2+} moments with S=7/2. Therefore, it is concluded that the moments of Eu^{2+} order ferromagnetically in $EuFe_2(As_{0.7}P_{0.3})_2$, in analogy with $EuFe_2P_2$[27]. Due to the proximity of superconducting transition and ferromagnetic ordering, the superconducting diamagnetic signal is hard to observe unless the applied magnetic field is very low. As shown in the inset of Fig. 2(c), ac magnetic susceptibility measured under μ_0H_{ac}=1 Oe shows a kink around 26 K. After subtraction of the paramagnetic contribution from Eu^{2+} moments, a clear diamagnetism is seen, confirming superconductivity in $EuFe_2(As_{0.7}P_{0.3})_2$.

The temperature dependence of resistivity under magnetic fields is shown in figure 3. With increasing magnetic fields, the resistive transition shifts towards lower temperature and becomes broadened, further affirming the superconducting transition. The $T_c(H)$, defined as a temperature where the resistivity falls to 50% of the normal state value, is plotted as a function of magnetic field in the inset of Fig. 3. The μ_0H_{c2}-T diagram shows a slight upward curvature, which is probably due to the multi-band effect[28]. The initial slope $\mu_0\partial H_{c2}/\partial T$ near T_c is -1.18 T/K, giving an upper critical field of $\mu_0H_{c2}(0)$ $\sim$ 30 T by linear extrapolation. It is also noted that the reentrant superconducting behavior is not obvious under magnetic field, in contrast with that in RNi$_2$B$_2$C$_2$ superconductors[25].

Figure 4 shows the specific heat (C) for the

$EuFe_2(As_{0.7}P_{0.3})_2$ sample. Two anomalies below 30 K are identified. One is a λ-shape peak with the onset at 20 K, indicating a second-order transition. With increasing magnetic fields, the anomaly shifts to higher temperatures and becomes broadened, in accordance with ferromagnetic nature of the transition. The other anomaly is much weaker but detectable at $\sim$26 K (shown in the upper-left panel of Fig. 4), which coincides with the superconducting transition.

To analyze the $C(T)$ data further, it is assumed that the total specific heat consists of the electronic, phonon and magnetic components. At high temperatures, the dominant contribution comes from the phonon component, which can be well described by the Debye model with only one adjustable parameter (i. e., Debye temperature Θ_D). The data fitting above 100 K gives Θ_D of 345 K. Since $\Theta_D\sim1/\sqrt{M}$ (M the molecular weight), the Θ_D for $EuFe_2(As_{0.7}P_{0.3})_2$ is calculated to be 348 K, in good agreement with the fitted value, using the Θ_D data of 380 K for $EuFe_2P_2$[27]. The upper bound of electronic specific heat coefficient is estimated to be 10 mJ/mol·K^2, which is comparable with those of other iron-arsenide superconductors. The electronic specific heat contribution is less than 2% of the total specific heat even at low temperatures, and thus is not taken into consideration in the following analysis.

By subtracting the lattice contribution, the specific anomaly due to the superconducting transition is more prominent. The jump in specific heat $\Delta C\approx300$ mJ/mol·K at T_c, which is of the same order as that in $Ba_{0.55}K_{0.45}Fe_2As_2$ crystals[30]. Meanwhile, the magnetic

entropy associated with the ferromagnetic transition is 16.5 J/mol·K, which amounts to 95% of $R\ln(2S+1)$ with $S=7/2$ for Eu^{2+} ions. The thermodynamic properties indicate that the superconducting transition and the ferromagnetic ordering are both of bulk nature.

A broad specific-heat hump below 90 K is evident after deduction of the lattice contribution, indicating existence of additional magnetic contribution. This anomaly is accompanied with the upward deviation from the linear temperature dependence in resistivity as shown in Fig. 1(a). Meanwhile, negative magnetoresistance was observed in the same temperature range, and reaches -6% at 30 K under $\mu_0 H=8$ T (data not shown here). All these features are probably attributed to the interaction between the moments of Eu^{2+} ions and conduction electrons. Such interaction may be responsible for the observed ferromagnetic ordering of Eu^{2+} moments below 20 K.

The isovalent substitution of As with P does not change the number of Fe $3d$ electrons, but generates chemical pressure, as manifested by the shrinkage of lattice. According to a coherent-incoherent scenario[23], the low energy physics of the FeAs-containing system is described by the interplay of the coherent excitations (associated with the itinerant carriers) and incoherent ones (modeled in terms of Fe localized magnetic moments). The internal chemical pressure generated via P doping results in the enhancement of coherent spectral weight, which weakens the SDW ordering and probably induces a magnetic quantum critical point (QCP). In the structural point of view, the P doping results in closer distance between As(P) and Fe planes. As a consequence, the low-energy band width becomes larger (correspondingly the coherent spectral weight is enhanced), according to the related band calculations[31]. Furthermore, the magnetic fluctuations near the QCP may induce superconductivity, as has been well documented in literatures.[32] This explains the simultaneous suppression of SDW transition and emergence of superconductivity in $EuFe_2(As_{0.7}P_{0.3})_2$ assuming that the QCP locates near P content of $\sim$30%.

The superconducting properties of $EuFe_2(As_{0.7}P_{0.3})_2$ bear similarity with those of $EuFe_2As_2$ crystals under hydrostatic pressure. As a matter of fact, the onset temperature of resistive drop is nearly the same in both cases, suggesting that the effect of internal chemical pressure is in analogy with application of external physical pressure. Thus it is expectable to find superconductivity in other iron arsenide systems via the P-doping strategy.

In summary, we have found superconductivity at 26 K in $EuFe_2(As_{0.7}P_{0.3})_2$. Moreover, ferromagnetic ordering of Eu^{2+} moments coexists with the superconductivity below 20 K. The observation of sizable anomalies in thermodynamic properties, concomitant with the transitions, indicates that both of them are bulk phenomena. The interplay of superconductivity and ferromagnetism may bring about exotic properties and provide clues to the superconductivity mechanism, which renders $EuFe_2(As_{1-x}P_x)_2$ worthy of further exploration.

[1] Y. Kamihara *et al.*, J. Am. Chem. Soc. **130**, 3296 (2008).
[2] X. H. Chen *et al.*, Nature **453**, 761 (2008); G. F. Chen *et al.*, Phys. Rev. Lett. **100**, 247002 (2008).
[3] Z. A. Ren *et al.*, Europhysics Lett. **83**, 17002 (2008); H. Kito, H. Eisaki and A. Iyo, J. Phys. Soc. Jpn. **77**, 063707 (2008).
[4] C. Wang *et al.*, Europhysics Lett. **83**, 67006 (2008).
[5] J. Dong *et al.*, Europhysics Lett. **83**, 27006 (2008).
[6] C. Cruz *et al.*, Nature **453**, 899 (2008).
[7] A. S. Sefat *et al.*, Phys. Rev. B **78**, 104505 (2008); G. H. Cao *et al.*, arXiv: 0807.1304 (2008).
[8] A. S. Sefat *et al.*, Phys. Rev. Lett. **101**, 117004 (2008); A. Leithe-Jasper, W. Schnelle, C. Geibel, and H. Rosner, Phys. Rev. Lett. **101**, 207004 (2008).
[9] G. H. Cao *et al.*, arXiv:0807.4328 (2008).
[10] J. L. Li *et al.*, New J. Phys. *in press.*
[11] H. H. Wen *et al.*, Europhysics Lett. **82**, 17009 (2008).
[12] M. Rotter, M. Tegel, and D. Johrendt, Phys. Rev. Lett. **101**, 107006 (2008).
[13] K. Sasmal *et al.*, Phys. Rev. Lett. **101**, 107007 (2008); G. F. Chen *et al.*, Chin. Phys. Lett. **25**, 3403 (2008).
[14] H. S. Jeevan *et al.*, Phys. Rev. B, **78**, 092406 (2008).
[15] M. S. Torikachvili, S. L. Budko, N. Ni, and P. C. Canfield, Phys. Rev. Lett. **101**, 057006 (2008); T. Park *et al.*, J. Phys.: Condensed Matter **20**, 322204 (2008).
[16] P. L. Alireza *et al.*, J. Phys.: Condensed Matter **21**, 012208 (2008).
[17] C. F. Miclea *et al.*, arXiv:0808.2026 (2008).
[18] R. Marchand, W. Jeitschko, J. Solid State Chem. **24**, 351(1978).
[19] Z. Ren *et al.*, Phys. Rev. B **78**, 052501 (2008).
[20] H. S. Jeevan *et al.*, Phys. Rev. B, **78**, 052502 (2008).
[21] S. Jiang *et al.*, New J. Phys. *in press.*
[22] Z. Ren *et al.*, arXiv:0810.2595 (2008).
[23] J. H. Dai *et al.*, arXiv:0808.0305 (2008).
[24] F. Izumi *et al.*, Mater. Sci. Forum. **198**, 321 (2000).
[25] H. Eisaki *et al.*, Phys. Rev. B (R), **50**, 647 (1994).
[26] T. M. McQueen *et al.*, Phys. Rev. B, **78**, 024521 (2008).
[27] H. Raffius *et al.*, J. Phys. Chem. Solids, **52**, 787 (1991).
[28] F. Hunte *et al.*, Nature **453**, 903 (2008).
[29] Anupam *et al.*, arXiv:0812.1131 (2008).
[30] N. Ni *et al.*, Phys. Rev. B, **78**, 014507 (2008).
[31] V. Vildosola, L. Pourovskii, R. Arita, S. Biermann, and A. Georges, Phys. Rev. B, **78**, 064518 (2008).
[32] P. Gegenwart, Q. Si, and F. Steglich, Nature Phys. **4**, 186 (2008).